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FILE 'REGISTRY	ENTERED	AΤ	09:24:31	ON	20	MAR	2003
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L1 STRUCTURE UPLOADED

L2 0 S L1 SSS SAM

L3 0 S L1 SSS FULL

FILE 'STNGUIDE' ENTERED AT 09:26:21 ON 20 MAR 2003

FILE 'REGISTRY' ENTERED AT 09:28:05 ON 20 MAR 2003

L4 STRUCTURE UPLOADED

L5 0 S L4 SSS SAM

L6 0 S L4 SSS FULL

FILE 'STNGUIDE' ENTERED AT 09:28:40 ON 20 MAR 2003

FILE 'STNGUIDE' ENTERED AT 09:39:06 ON 20 MAR 2003

FILE 'REGISTRY' ENTERED AT 09:41:32 ON 20 MAR 2003

L7 STRUCTURE UPLOADED

L8 0 S L7 SSS SAM

L9 1 S L7 SSS FULL

FILE 'CAPLUS' ENTERED AT 09:42:11 ON 20 MAR 2003

L10 3 S L9

L11 3 S L10 AND (CARCINOM? OR SARCOM? OR LYMPHOM? OR LEUKEM? OR VIRAL

FILE 'STNGUIDE' ENTERED AT 09:44:54 ON 20 MAR 2003

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09/676,034

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L1 STRUCTURE UPLOADED

=> d 11

L1 HAS NO ANSWERS

L1 STR

G1

G2 O, N

Structure attributes must be viewed using STN Express query preparation.

=> s l1 sss full

FULL SEARCH INITIATED 21:22:40 FILE 'REGISTRY'
FULL SCREEN SEARCH COMPLETED - 18 TO ITERATE

100.0% PROCESSED

18 ITERATIONS

2 ANSWERS

SEARCH TIME: 00.00.02

L2

2 SEA SSS FUL L1

=> file caplus

=> s 12

L3 1 L2

=> d 13 abs ibib hitstr 1

L3 ANSWER 1 OF 1 CAPLUS COPYRIGHT 2001 ACS GI

$$R^{1}$$
 R^{2}
 $A-E-Y$
 R^{3}
 X

AB The title compds. [I; R1 = (un)substituted aryl, cyclo(lower)alkyl, heterocyclyl; R2 = H, lower alkyl, etc.; R3 = H, halo, OH, etc.; A = a single bond, O, NH; E = lower alkylene, lower alkenylene, etc.; X = CH:CH, CH:N, S; Y = (un)substituted aryl, condensed heterocyclyl, etc.] and their pharmaceutically acceptable salts, useful in treatment and/or prevention of hypertension, heart failure, renal insufficiency, edema, ascites, vasopressin parasecretion syndrome, hepatocirrhosis, hyponatremia, hypokalemia, diabetic, circulation disorder, cerebrovascular disease, Meniere's disease or motion sickness, were prepd. Thus, the title compd.

II showed IC50 of 1.5 nM against vasopressin 1 receptor binding.

ACCESSION NUMBER:

1998:394328 CAPLUS

DOCUMENT NUMBER:

129:67773

TITLE:

Preparation of benzamide derivatives having a

vasopressin antagonistic activity

INVENTOR(S):

Setoi, Hiroyuki; Ohkawa, Takehiko; Zenkoh, Tatsuya;

Sawada, Hitoshi; Sawada, Yuki; Oku, Teruo

PATENT ASSIGNEE(S):

Fujisawa Pharmaceutical Co., Ltd., Japan; Setoi,

Hiroyuki; Ohkawa, Takehiko; Zenkoh, Tatsuya; Sawada, Hitoshi; Sawada, Yuki; Oku, Teruo

SOURCE:

PCT Int. Appl., 332 pp. CODEN: PIXXD2

DOCUMENT TYPE:

Patent

LANGUAGE:

English

FAMILY ACC. NUM. COUNT:

PATENT INFORMATION:

PA'	rent	NO.		KI	ND	DATE		APPLICATION NO.				ο.	DATE					
WO	9824			A	1	1998	0611		W	0 19	 97-J	- P419	 2	1997	 1118			
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								1	WO 1	997-	JP419	92	W	1997	1118			

OTHER SOURCE(S):

MARPAT 129:67773

IT208767-99-1P 208768-81-4P

RL: BAC (Biological activity or effector, except adverse); SPN (Synthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses)

(prepn. of benzamide derivs. having a vasopressin antagonistic activity)

208767-99-1 CAPLUS RN

CN Carbamic acid, [4-[[[2-methoxy-4-[[methyl[4-methyl-2-[[6-(4-methyl-1-methyl-2-[[6-(4-methyl-1-methyl-2-[[6-(4-methyl-1-methyl-2-[[6-(4-methyl-1-methyl-2-[[6-(4-methyl-1-methyl-2-[[6-(4-methyl-1-methyl-2-[[6-(4-methyl-1-methyl-2-[[6-(4-methyl-1-methyl-1-methyl-2-[[6-(4-methyl-1-methyl-1-methyl-2-[[6-(4-methyl-1-methyl-1-methyl-1-methyl-2-[[6-(4-methyl-1-methyl-1-methyl-1-methyl-1-methyl-1-methyl-1-methyl-1-methyl-1-methyl-1-methyl-1-methyl-1-methyl-1-methyl-1-methyl-1-methyl-1-methyl-1-methyl-1-methyl-1-methyl-1-methyl-1-methyl-1-methyl-1-methyl-1-methyl-1-methyl-1-methyl-1-methyl-1-methyl-1-methyl-1-methyl-1-methyl-1-methyl-1-methyl-1-methyl-1-methyl-1-methyl-1-methyl-1-methyl-1-methyl-1-methyl-1-methyl-1-methyl-1-methyl-1-methyl-1-methyl-1-methyl-1-methyl-1-methyl-1-methyl-1-methyl-1-methyl-1-methyl-1-methyl-1-methyl-1-methyl-1-methyl-1-methyl-1-methyl-1-methyl-1-methyl-1-methyl-1-methyl-1-methyl-1-methyl-1-methyl-1-methyl-1-methyl-1-methyl-1-methyl-1-methyl-1-methyl-1-methyl-1-methyl-1-methyl-1-methyl-1-methyl-1-methyl-1-methyl-1-methyl-1-methyl-1-methyl-1-methyl-1-methyl-1-methyl-1-methyl-1-methyl-1-methyl-1-methyl-1-methyl-1-methyl-1-methyl-1-methyl-1-methyl-1-methyl-1-methyl-1-methyl-1-methyl-1-methyl-1-methyl-1-methyl-1-methyl-1-methyl-1-methyl-1-methyl-1-methyl-1-methyl-1-methyl-1-methyl-1-methyl-1-methyl-1-methyl-1-methyl-1-methyl-1-methyl-1-methyl-1-methyl-1-methyl-1-methyl-1-methyl-1-methyl-1-methyl-1-methyl-1-methyl-1-methyl-1-methyl-1-methyl-1-methyl-1-methyl-1-methyl-1-methyl-1-methyl-1-methyl-1-methyl-1-methyl-1-methyl-1-methyl-1-methyl-1-methyl-1-methyl-1-methyl-1-methyl-1-methyl-1-methyl-1-methyl-1-methyl-1-methyl-1-methyl-1-methyl-1-methyl-1-methyl-1-methyl-1-methyl-1-methyl-1-methyl-1-methyl-1-methyl-1-methyl-1-methyl-1-methyl-1-methyl-1-methyl-1-methyl-1-methyl-1-methyl-1-methyl-1-methyl-1-methyl-1-methyl-1-methyl-1-methyl-1-methyl-1-methyl-1-methyl-1-methyl-1-methyl-1-methyl-1-methyl-1-methyl-1-methyl-1-methyl-1-methyl-1-methyl-1-methyl-1-methyl-1-methyl-1-methyl-1-methyl-1-methyl-1-methylpiperazinyl)-6-oxohexyl]oxy]phenyl]amino]carbonyl]phenyl]amino]carbonyl]-1H-benzimidazol-2-yl]-, methyl ester (9CI) (CA INDEX NAME)

PAGE 2-A

RN 208768-81-4 CAPLUS

CN Carbamic acid, [4-[[[2-methoxy-4-[[methyl[4-methyl-2-[[6-(4-methyl-1-piperazinyl)-6-oxohexyl]oxy]phenyl]amino]carbonyl]phenyl]amino]carbonyl]-1H-benzimidazol-2-yl]-, methyl ester, dihydrochloride (9CI) (CA INDEX NAME)

PAGE 1-A

PAGE 2-A

H
NH-C-OMe

●2 HCl

=> file reg

09/676,034

Uploading 0344.str

STRUCTURE UPLOADED

=> d 14

L4 HAS NO ANSWERS

L4

G1

G2 O, N

G3 O, N

Structure attributes must be viewed using STN Express query preparation.

0 ANSWERS

=> s 14 sss full

FULL SEARCH INITIATED 21:24:17 FILE 'REGISTRY' FULL SCREEN SEARCH COMPLETED - 105 TO ITERATE

100.0% PROCESSED 105 ITERATIONS

SEARCH TIME: 00.00.02

L5 0 SEA SSS FUL L4